



Australian Government



PM_{2.5} Sources at Mayfield, Newcastle

Eduard Stelcer, David D. Cohen, Armand J. Atanacio

**Australian Nuclear Science and Technology Organisation
Institute for Environmental Research, Locked Bag 2001, Kirrawee NSW**

Introduction

Long term PM_{2.5} sampling at Mayfield sampling site (Feb1998 – Dec2013)

Ion Beam Analysis (IBA) of collected samples

Positive Matrix Factorisation (PMF)



Mayfield



Mayfield

**Inner city suburb of Newcastle,
approximately 150 km north of Sydney**

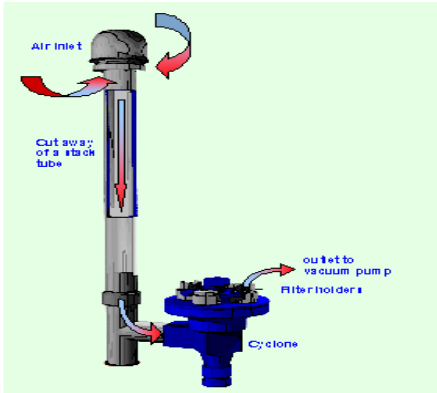
**It is close to highly industrialised areas and
the port of Newcastle**

**The region was originally developed as an
area for heavy industry, specifically iron and
steel making**

Sampling

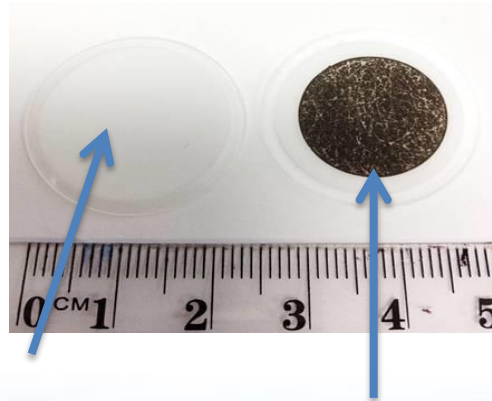
Every Wednesday and Sunday from February 1998 to December 2013

**Standard IMPROVE PM_{2.5}
cyclone sampling unit**



22 l/min flow rate

Stretched Teflon filters



**Clean
filter**

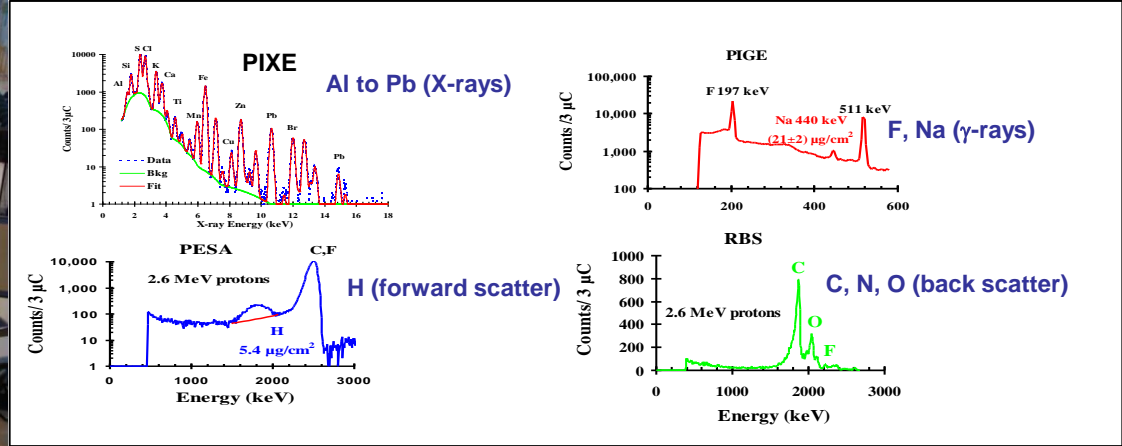
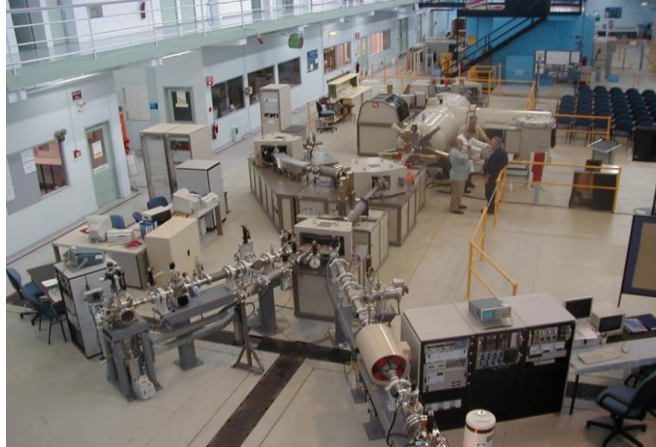
**Exposed
filter**

1538 days sampled

1421 good sampling days

Analysis

Accelerator based Ion Beam Analysis (IBA) techniques

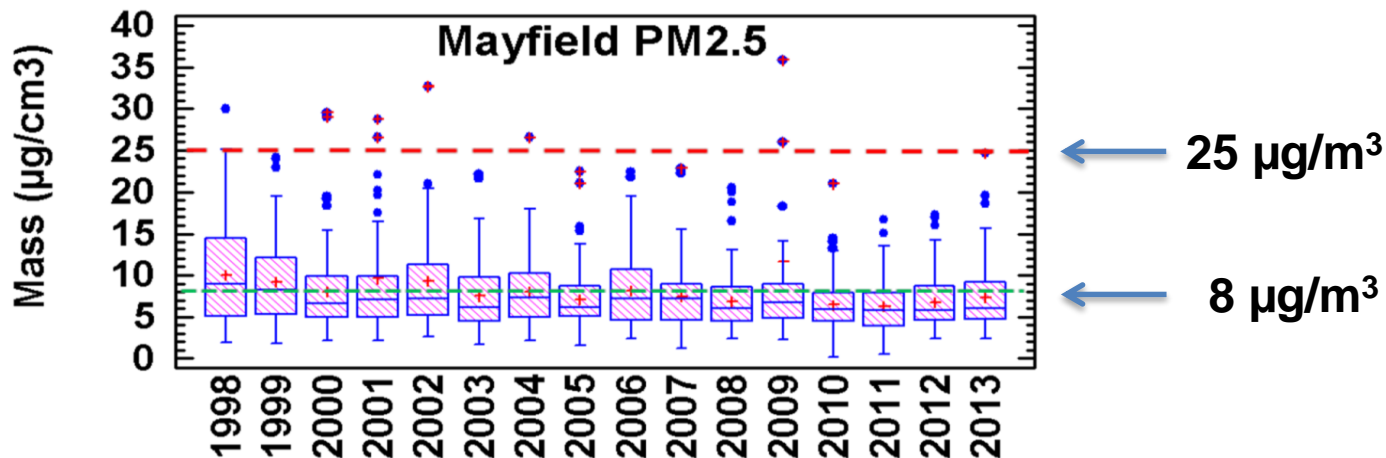


H, Na, Al, Si, P, S, Cl, K, Ca, Ti, V, Cr, Mn, Fe, Co, Ni, Cu, Zn, Se, Br, & Pb

Laser Integrating Plate Method (LIMP)

BC

Gravimetric Mass and the Australian NEPC Goals



Box and whisker plot of the 1,421 fine mass measurements at Mayfield site (+) symbols are annual average values

horizontal bars in the boxes are annual median values

The **average gravimetric mass** over the study period was **$(8.11 \pm 12) \mu\text{g}/\text{m}^3$**

The **maximum mass** value of **$402 \mu\text{g}/\text{m}^3$** occurred during significant desert dust storms in September 2009

Composition of the fine particles at Mayfield (1998-2013)

PM _{2.5}	Av.	Median	SD	Max	MDL
H (ng m ⁻³)	271	180	1,118	41,849	4.2
Na (ng m ⁻³)	404	307	427	3,962	58
Al (ng m ⁻³)	44	26	123	4,107	3.2
Si (ng m ⁻³)	106	66	344	11,899	1.6
P (ng m ⁻³)	3.1	1.5	5.0	98	1.7
S (ng m ⁻³)	414	339	285	2,612	1.4
Cl (ng m ⁻³)	431	272	492	4,288	1.5
K (ng m ⁻³)	63	43	88	2,377	0.8
Ca (ng m ⁻³)	49	33	64	1,511	0.9
Ti (ng m ⁻³)	4.9	2.7	26	987	0.6
V (ng m ⁻³)	2.6	1.6	3.2	41	0.6
Cr (ng m ⁻³)	0.6	0.3	0.9	9	0.5
Mn (ng m ⁻³)	89	7	180	1,633	0.5
Fe (ng m ⁻³)	135	57	354	10,704	0.6
Co (ng m ⁻³)	0.6	0.2	2.2	70	1.0
Ni (ng m ⁻³)	1.4	0.9	4.8	155	0.5
Cu (ng m ⁻³)	2.3	1.5	3.0	51	0.6
Zn (ng m ⁻³)	34	13	83	1,483	0.7
Se (ng m ⁻³)	0.7	0.5	0.9	9	1.5
Br (ng m ⁻³)	4.7	3.1	5.3	61	1.9
Pb (ng m ⁻³)	15	5.9	27	336	3.7
BC (ng m ⁻³)	1,237	1,048	813	7,367	29
Salt (µg m ⁻³)	1.03	0.78	1.1	10	0.15
Soil (µg m ⁻³)	0.78	0.47	2.0	69	0.02
Organics (µg m ⁻³)	1.87	0.89	12	460	0.05
Ammonium	1.71	1.40	1.2	11	0.06
Sulfate (µg m ⁻³)	6.59	5.35	14	519	0.04
RCM (µg m ⁻³)	81	80	12	137	
RCM%	8.11	6.62	12	402	0.16
Mass (µg m ⁻³)					

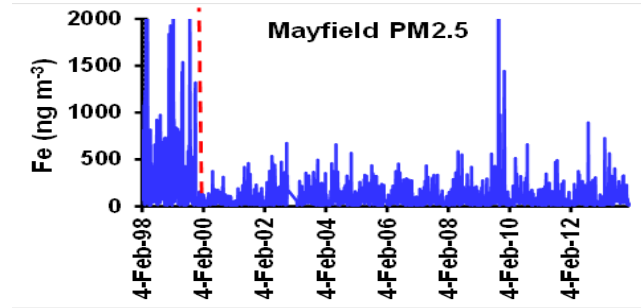
Salt = 2.54Na

Soil = 2.2Al + 2.49Si + 1.63Ca + 1.94Ti + 2.42Fe

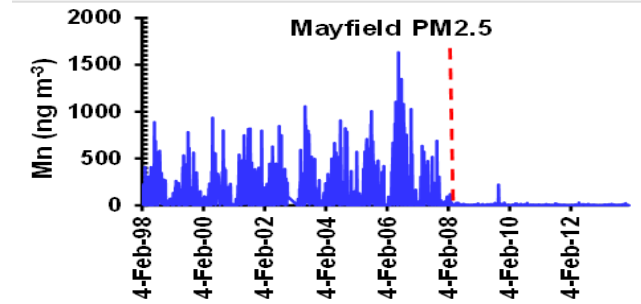
Organics = 11(H - 0.25S)

Ammonium Sulfate = 4.125S

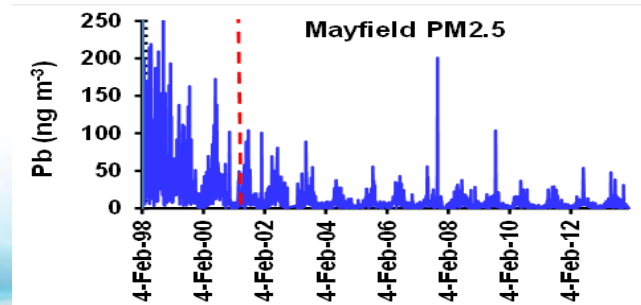
Time Series Plots (1998 – 2013)



October 1999
Fe – dropping
from over 500 ng/m³ to less than 300 ng/m³

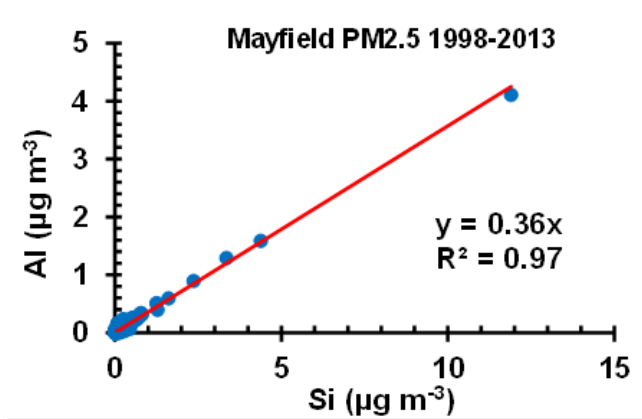


March 2008
Mn – dropping
from over 500 ng/m³ to almost 0 ng/m³

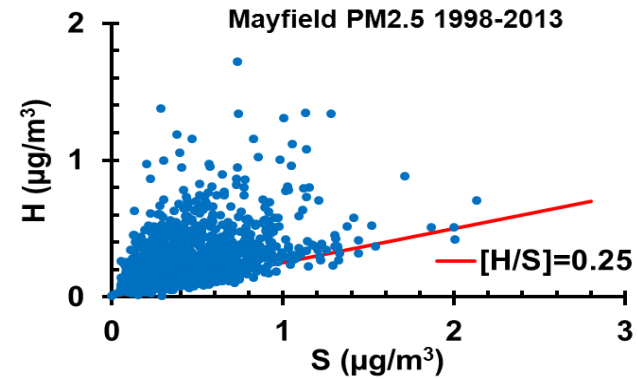


January 2001
Pb – dropping
from over 50 ng/m³ to less than 10 ng/m³

Scatter Plots



[Al/Si] = 0.36
Clay Alumina-Silicates (Soil)



[H/S] = 0.25
Ammonium sulfate [(NH₄)₂SO₄]

Positive Matrix Factorisation (PMF2)

Each sampling site has its own fine particle sources and each source has its own elemental fingerprint

PMF is a one-step **receptor** modelling technique for determining the elemental **fingerprints** for key pollution sources and their **contributions** to total pollution

PMF does not automatically assign a source name to each fingerprint - it only provides a source factor which is then given a name by the user

This is based on sampling site knowledge and experience in identifying the source associated with elemental fingerprint of each factor

PMF source fingerprints

Al, Si, K, Ca, Ti, Fe
[Al/Si] = 0.37

H, Na, S

Na, S, Cl, K
[Cl/Na] = 1.47

H, S, K, BC

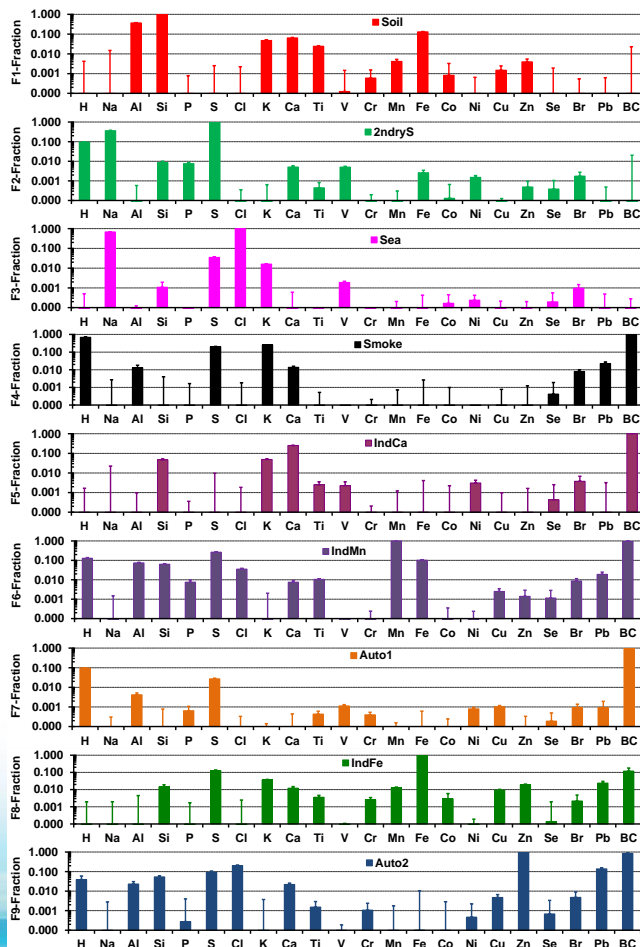
Si, K, Ca, BC

H, S, Mn, Fe, BC

H, S, Bc

S, Fe, BC

H, Si, S, Cl, Zn,
Pb, BC



Percentages of total elemental concentrations in sources

74% Al, 89% Si, 61% Ti

11% H, 41% Na, 78% P,
78% S, 58% V, 44% Ni

59% Na, 99% Cl

66% H, 84% K, 51% Br,
62% Pb, 28% BC

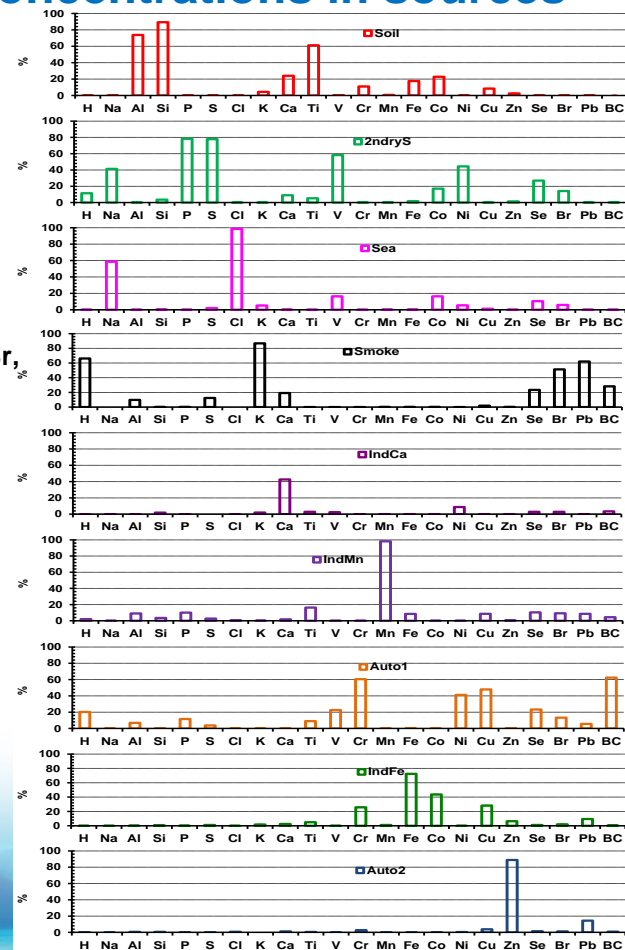
43% Ca

98% Mn

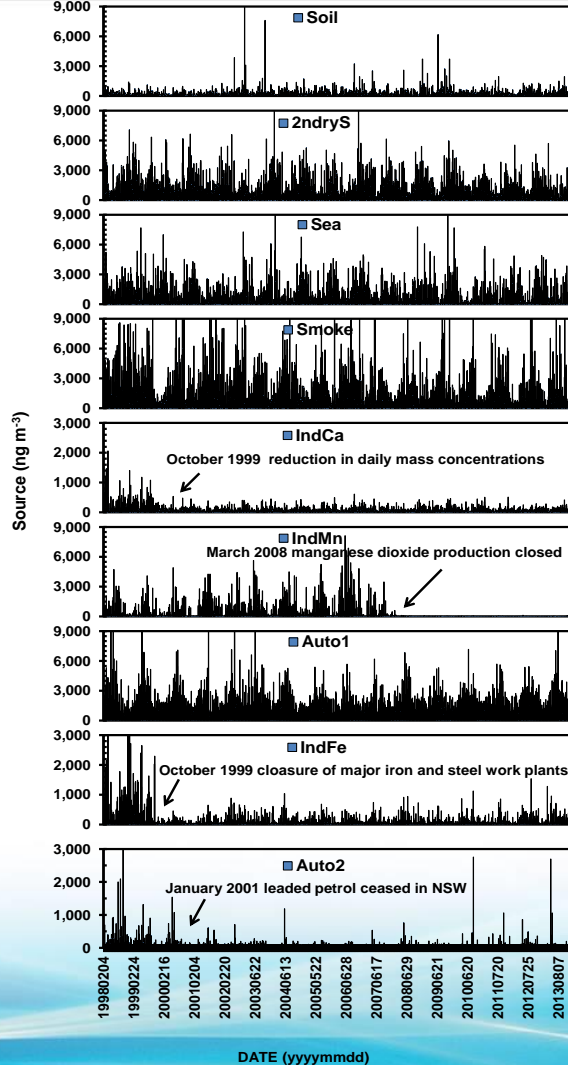
60% Cr, 41% Ni,
48% Cu, 57% BC

72% Fe

89% Zn



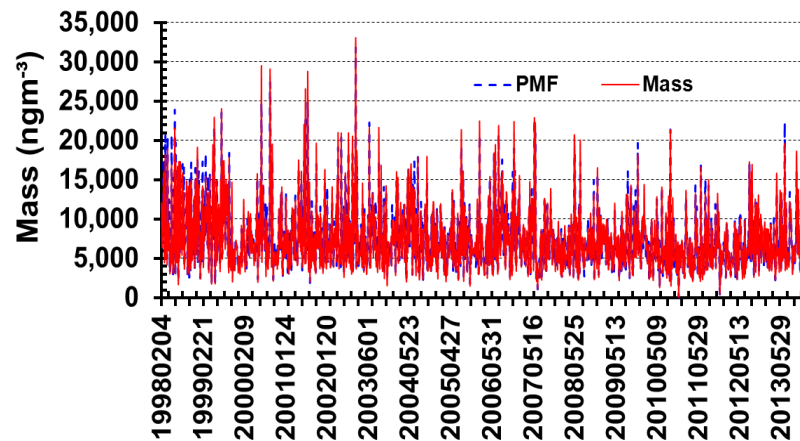
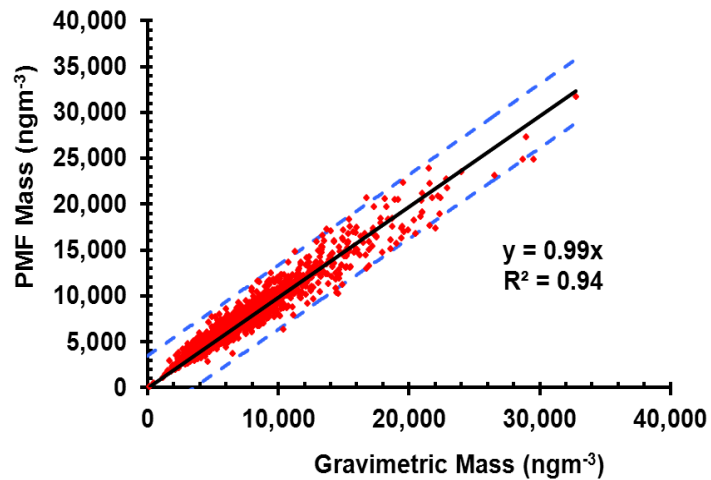
Daily source concentrations



PMF

Average contributions to the total PM_{2.5} mass 1998 - 2013

Factor	Fingerprint	%Mass
F1	Soil	4.6±0.25
F2	2ndryS	19.7±0.42
F3	Sea	17.3±0.37
F4	Smoke	23.0±0.35
F5	IndCa	1.72±0.51
F6	IndMn	4.56±0.15
F7	Auto1	26.1±0.48
F8	IndFe	2.08±0.27
F9	Auto2	0.97±0.13



Conclusion

IBA is a powerful non-destructive multi-elemental analytical tool which can provide information on elemental concentrations for most of the elements that can be found in atmospheric particles.

The combination of long-term IBA data sets with PMF can be used to determine source fingerprints and their contribution to total pollution on a local scale.

This source information can be more meaningful to local councils and regulatory bodies than elemental concentration alone.



Thank you for your attention